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DATE: Monday, April 24, 2006

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	DB=PC	GPB,USPT; PLUR=YES; OP=OR	
	L10	L9 and ((800/278).ccls. or (800/294).ccls. or (435/468).ccls. or (435/469).ccls. or (800/290).ccls.)	43
	L9	17 not 15	83
	L8	select\$ and marker	153392
	L7	L6 same (select\$ or marker)	84
	L6	indoleacetamide or (indole adj acetamide) or naphthaleneacetamide or (naphthalene adj acetamide)	248
	L5	L4 and ((800/278).ccls. or (800/294).ccls. or (435/468).ccls. or (435/469).ccls. or (800/290).ccls.)	8
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	L3	ebinuma.in. and hiroyasu.in.	10
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	L1	etsuko.in. and matsunaga.in.	7

**END OF SEARCH HISTORY** 

## Hit List

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**Search Results -** Record(s) 1 through 8 of 8 returned.

☐ 1. Document ID: US 20040163143 A1

Using default format because multiple data bases are involved.

L5: Entry 1 of 8

File: PGPB

Aug 19, 2004

PGPUB-DOCUMENT-NUMBER: 20040163143

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040163143 A1

inte t

TITLE: Method for efficiently producing transgenic plant using auxin precursor

PUBLICATION-DATE: August 19, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY Matsunaga, Etsuko Tokyo JΡ Sugita, Koichi Tokyo JP Ebinuma, Hiroyasu Tokyo JP

US-CL-CURRENT: 800/278

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KođC	Draw, De
							-					

☐ 2. Document ID: US 20030221210 A1

L5: Entry 2 of 8

File: PGPB

Nov 27, 2003

PGPUB-DOCUMENT-NUMBER: 20030221210

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030221210 A1

09/850,09/ ABN ×18/01

TITLE: Method for introducing a gene into a plant using and adventitious bud redifferentiation gene under the control of a light-inducible promoter as a selectable marker gene, and vector for introducing a gene into a plant using the

PUBLICATION-DATE: November 27, 2003

INVENTOR-INFORMATION:

CON 09/354,305 7/16/99 COUNTRY 6,294,714 NAME CITY STATE Matsunaga, Etsuko Tokyo JP

Kasahara, Takehide Tokyo JP Sugita, Koichi

Tokyo

JР

Ebinuma, Hiroyasu

Tokyo

JP

US-CL-CURRENT: 800/278; 435/320.1

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims KWC Draw. D

☐ 3. Document ID: US 20030033639 A1

L5: Entry 3 of 8

File: PGPB

Feb 13, 2003

PGPUB-DOCUMENT-NUMBER: 20030033639

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030033639 A1

10/201, 110 fled 7/24/2

TITLE: Method for introducing gene into plant having improved transformation

efficiency

PUBLICATION-DATE: February 13, 2003

Fral R

INVENTOR - INFORMATION:

NAME

CITY STATE COUNTRY

Matsunaga, EtsukoTokyoJPSugita, KoichiTokyoJPEbinuma, HiroyasuTokyoJP

US-CL-CURRENT: 800/294

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims KMC Draw De

☐ 4. Document ID: US 20020123623 A1

L5: Entry 4 of 8

File: PGPB

Sep 5, 2002

PGPUB-DOCUMENT-NUMBER: 20020123623

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020123623 A1

NO

TITLE: Transcription factor controlling phenylpropanoid biosynthesis pathway

PUBLICATION-DATE: September 5, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY

Kawaoka, Akiyoshi Tokyo JP <u>Ebinuma, Hiroyasu</u> Tokyo JP

US-CL-CURRENT: 536/23.6; 435/320.1, 530/370, 800/278

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims KMC Draw, De

☐ 5. Document ID: US 6767735 B1

L5: Entry 5 of 8

File: USPT

Jul 27, 2004

US-PAT-NO: 6767735

DOCUMENT-IDENTIFIER: US 6767735 B1

N

TITLE: Vector for introducing a gene into a plant using a selectable marker

Full Title Cit	tation Front Review	Classification Date	Reference Sequences	Attachments Claims RMC Dr	avu De

☐ 6. Document ID: US 6326192 B1

L5: Entry 6 of 8

File: USPT

Dec 4, 2001

US-PAT-NO: 6326192

DOCUMENT-IDENTIFIER: US 6326192 B1

TITLE: Vector for gene transfer into plant allowing optional deletion of marker

gene

Full Title Citation Front Review Classification Date Reference **Sequences Attachments** Claims KWIC Draw Do

☐ 7. Document ID: US 6294714 B1

L5: Entry 7 of 8

File: USPT

Sep 25, 2001

US-PAT-NO: 6294714

DOCUMENT-IDENTIFIER: US 6294714 B1

TITLE: Method for introducing a gene into a plant using an adventitious bud redifferentiation gene under the control of a light-inducible promoter as a selectable marker gene, and vector for introducing a gene into a plant using the same

Full Title Citation Front Review Classification Date Reference Seguences Attachments Claims KMC Draw. De

□ 8. Document ID: US 5965791 A

L5: Entry 8 of 8

File: USPT

Oct 12, 1999

US-PAT-NO: 5965791

DOCUMENT-IDENTIFIER: US 5965791 A

V

TITLE: Vector for introducing a gene into a plant, and methods for producing transgenic plants and multitudinously introducing genes into a plant using the vector

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims KMC Draw De

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	Terms	Documents			
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£10: Entry 29 of 43

File: USPT

Jun 22, 2004

DOCUMENT-IDENTIFIER: US 6753459 B2

\*\* See image for Certificate of Correction \*\*

TITLE: Transgenic plants and methods for production thereof

Current US Original Classification (1): 800/278

<u>Current US Cross Reference Classification</u> (11): 800/294

## CLAIMS:

- 1. A genetic construct, comprising: a conditionally lethal first gene expressible in a plant cell of a plant, said conditionally lethal first gene being a gene encoding indoleacetamide hydrolase (IAMH); and a second gene expressible in said plant cell, said second gene, when expressed in said plant cell, conferring a non-naturally occurring trait of interest on said plant cell, said second gene being selected from the group consisting of: (a) a gene which, when expressed in said plant cell, confers insect resistance on said plant cell; (b) a gene which, when expressed in said plant cell, confers an output trait on said plant cell; (c) a gene encoding an industrially useful enzyme; (d) a gene encoding a pharmaceutically active compound; (e) a gene encoding rennin or hirudin; and (f) a gene encoding an antisense RNA.
- 15. A transgenic plant, comprising: a conditionally lethal first gene expressible in a plant cell of said transgenic plant, said conditionally lethal first gene being a gene encoding indoleacetamide hydrolase (IAMH); and a second gene expressible in said plant cell of said transgenic plant, said second gene, when expressed in said plant cell, conferring a non-naturally occurring trait of interest on said plant cell, said second gene being selected from the group consisting of: (a) a gene which, when expressed in said plant cell, confers insect resistance on said plant cell; (b) a gene which, when expressed in said plant cell, confers an output trait on said plant cell; (c) a gene encoding an industrially useful enzyme; (d) a gene encoding a pharmaceutically active compound; (e) a gene encoding rennin or hirudin; and (f) a gene encoding an antisense RNA.
- 16. A method for <u>selectively</u> removing at least one plant from a growing environment, comprising: transforming at least one plant cell with a genetic construct including: a conditionally lethal first gene expressible in said at least one plant cell, said conditionally lethal first gene being a gene encoding <u>indoleacetamide</u> hydrolase (IAMH); and a second gene expressible in said at least one plant cell, said second gene, when expressed in said at least one plant cell, conferring a non-naturally occurring trait of interest on said at least one plant cell, said second gene being <u>selected</u> from the group consisting of: (a) a gene which, when expressed in said plant cell, confers insect resistance on said plant cell; (b) a gene which, when expressed in said plant cell, confers an output trait on said plant cell; (c) a gene encoding an industrially useful enzyme; (d) a gene encoding a pharmaceutically active compound; (e) a gene encoding rennin or hirudin; and (f) a gene encoding an antisense RNA.; regenerating the at least one plant cell to at least one whole plant; and applying a chemical agent to said at least one

whole plant, said chemical agent being configured to be converted into a phytotoxic agent of said at least one whole plant by one or more gene products of said conditionally lethal gene, wherein said chemical agent comprises an indoleamide or a related auxin derivative that is a substrate for IAMH.

20. A method for <u>selecting</u> a germinating seed or plant embryo comprising a transgene, comprising: providing at least one transgenic plant cell of a plant seed or plant embryo, said at least one transgenic plant cell including a transgene encoding <u>indoleacetamide</u> hydrolase (IAMH); culturing the at least one transgenic plant cell on a medium comprising an auxin transport inhibitor and an indoleamide or a related auxin derivative that is a substrate for IAMH; and visually identifying the at least one transgenic plant cell by its expression of a sublethal auxin-overproduction phenotype.

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L10: Entry 12 of 43

File: PGPB

Dec 2, 2004

DOCUMENT-IDENTIFIER: US 20040244076 A1

TITLE: Method for large scale mutagenesis in crop plants

Current US Classification, US Primary Class/Subclass: 800/278

Current US Classification, US Secondary Class/Subclass: 435/468

Detail Description Paragraph:

[0094] (3) Selection markers and GUS reporter. In addition to the kanamycin selection needed for transformation and the GUS reporter utilized in the trapping systems, a number of markers were used to select for transposition events (Fedoroff and Smith, 1993; Sundaresan et al., 1995). To that end, sterilized seeds were germinated and grown in 0.8% agar-containing Nitsh mediun supplemented with either one or a combination of the following compounds: 20 .mu.g/ml hygromycin (Calbiochem); 0.25 .mu.g/ml naphthalene acetamide (NAM, Sigma); and 100 p.p.b. or 2 p.m. chlorosulfuron (DuPont). GUS staining was done according to Jefferson (1987) and tissue clearing was done according to Beeckman and Engler (1994).

Detail Description Paragraph:

[0099] The indole acetic hydrolase (iaaH) gene confers sensitivity to NAM. Sensitive plants develop a callus-like tissue at the root base and die about three weeks after germination, as shown in FIG. 4C. Plants, transformed with Bam35S-Ac are sensitive to 0.25 .mu.g/ml naphthalene acetamide (FIG. 4C, left) while the wild-type is resistant (FIG. 4C, right). NAM sensitivity can be used as a negative selection marker to select against Bam35S-Ac, thus stabilizing new insertions, and/or to select against the donor site in DsE and DsG.

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